

High CO_x Hydrogenation to Methane over Co_aCe_{1-a}O_x at Low Temperature Prepared by Co-precipitation Method

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Conversion of CO₂ to methane, which is a versatile feedstock, is one of the most effective solutions to solve the global warming problem by increasing CO₂ concentration. Noble metal-based catalysts are believed to be more active in CO₂ methanation, however, those catalysts are limited in industrial applications due to their higher volatility, higher cost than other catalysts as well as easy sintering at relatively high temperature. Therefore, CO₂ methanation via heterogeneous catalysts has recently attracted a considerable amount of attention.

Cobalt-Ceria catalysts, Co_aCe_{1-a}O_x, with various Co²⁺/Ce⁴⁺ molar ratios from 3:7 to 9:1 were prepared by co-precipitation method and calcined at 500°C for 5 h in the air stream, then were subsequently characterized by means of X-ray diffraction (XRD), N₂ physisorption and desorption and H₂-Temperature programmed reduction (H₂-TPR). The effects of CeO₂ content on catalytic performance were evaluated in the separated CO and CO₂ methanation reaction in the fixed-bed reactor by the excess of H₂ at the temperature range of 120°C–350°C. The calcination temperature effects were also investigated at 350°C and 500°C of Co₃O₄ and Co_{0.9}Ce_{0.1}O_x catalysts.